

GenCore version 4.5
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OM protein - protein search, using sw model
Run on: April 25, 2000, 20:10:14 ; Search time 85.2 Seconds
(without alignments)
176.812 Million cell updates/sec

Title: US-09-125-005-6
Sequence: 1 MAQSTATSPDGTTTEBHLWS.....PDCKARKQPIKEEFEAETH 636
23688106 residues

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 188963 seqs,

Total number of hits satisfying chosen parameters: 188963

Minimum DB seq length: 0

Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database : A_Geneseq_36:*

Pre^d. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	3384	100.0	636	1 W36184	Human p53 tumour s
2	3367.5	99.5	635	1 W30661	Human NBS-1 alpha
3	3304.5	97.7	637	1 W36182	Monkey p53 tumour
4	3130	92.5	588	1 W36189	Human p53 tumour s
5	3058	90.4	587	1 W36187	Human p53 tumour s
6	2830.5	83.6	589	1 W36185	Mouse p53 tumour s
7	2624	77.5	499	1 W36190	Human p53 tumour s
8	2574	76.1	499	1 W36183	Monkey p53 tumour s
9	2330.5	68.9	506	1 W36188	Human p53 tumour s
10	726.5	21.5	401	1 W28487	Human p53 protein
11	725.5	21.4	401	1 W28488	Human p53 protein
12	724.5	21.4	355	1 W13950	De1356-333 modified
13	724.5	21.4	363	1 W13954	Modified p53 varia
14	724.5	21.4	393	1 R22238	Sequence of 53 KD
15	724.5	21.4	393	1 R26758	P53. Synthetic onc
16	724.5	21.4	393	1 R79658	Human p53 protein
17	724.5	21.4	393	1 R94623	Human p53 protein. Recom
18	724.5	21.4	393	1 R91933	Wild type p53 prot
19	724.5	21.4	393	1 W02617	Human p53 tumour s
20	724.5	21.4	393	1 W05344	Human p53. New hum
21	724.5	21.4	393	1 W13948	Human wild-type p5
22	724.5	21.4	393	1 W57242	Human p53 protein
23	724.5	21.4	393	1 W57243	Human p53 protein
24	724.5	21.4	393	1 W48658	Amino acid sequence
25	724.5	21.4	393	1 W69217	Human wild-type p5
26	724.5	21.4	393	1 W69718	Human p53 used in
27	724.5	21.4	393	1 Y03191	Amino acid sequence
28	724.5	21.4	438	1 R74277	Tumour suppressor
29	724.5	21.4	533	1 W19763	p53 GM-CSF immuno
30	723.5	21.4	363	1 W13971	Modified p53 varia
31	723.5	21.4	393	1 W13949	T284R modified hum
32	723.5	21.4	393	1 W13953	T284K modified hum
33	723.5	21.4	393	1 W57244	Human p53 protein
34	723.5	21.4	393	1 W57245	Human p53 protein

ALIGNMENTS

Human p53 protein.	35	723.5	21.4	393	1 W84270
p53 tumour suppressor	36	723.5	21.4	438	1 W50088
Modified p53 varia	37	720.5	21.3	363	1 W13972
Human p53 mutant R	38	720.5	21.3	393	1 W05347
Human tumour-deriv	39	720.5	21.3	393	1 W13551
Human p53 amino ac	40	719.0	21.3	354	1 R51874
Modified p53 varia	41	719.5	21.3	363	1 W13973
Modified p53 varia	42	719.5	21.3	363	1 W13974
Human p53 mutant N	43	719.5	21.3	393	1 W05345
Human p53 mutant R	44	719.5	21.3	393	1 W03346
Modified p53 varia	45	719.5	21.3	393	1 W13968

RESULT 1	W36184	Standard; Protein; 636 AA.	AC W36184;	AC W36184;	AC W36184;
ID W36184	W36184		ID W36184;	ID W36184;	ID W36184;
DE			DE 27-APR-1998 (first entry)	DE 27-APR-1998 (first entry)	DE 27-APR-1998 (first entry)
KW Human p53 tumour suppressor-related protein SR-p70a.			KW SR-p70a; human; transcription factor; p53; tumour suppressor gene; homology; differential splicing; diagnosis; cancer; neuroblastoma; gene therapy; apoptosis.	KW SR-p70a; human; transcription factor; p53; tumour suppressor gene; homology; differential splicing; diagnosis; cancer; neuroblastoma; gene therapy; apoptosis.	KW SR-p70a; human; transcription factor; p53; tumour suppressor gene; homology; differential splicing; diagnosis; cancer; neuroblastoma; gene therapy; apoptosis.
OS Homo sapiens.			OS Homo sapiens.	OS Homo sapiens.	OS Homo sapiens.
PN W0972836-Al.			PN W0972836-Al.	PN W0972836-Al.	PN W0972836-Al.
PD 07-AUG-1997.			PD 07-AUG-1997.	PD 07-AUG-1997.	PD 07-AUG-1997.
PR 03-FEB-1997; F00214.			PR 03-FEB-1997; F00214.	PR 03-FEB-1997; F00214.	PR 03-FEB-1997; F00214.
PA (SIFI) SANOFI SA.			PA (SIFI) SANOFI SA.	PA (SIFI) SANOFI SA.	PA (SIFI) SANOFI SA.
PI Caput D, Ferrara P, Kaghad AM;			PI Caput D, Ferrara P, Kaghad AM;	PI Caput D, Ferrara P, Kaghad AM;	PI Caput D, Ferrara P, Kaghad AM;
DR W71; 97-402550/37.			DR W71; 97-402550/37.	DR W71; 97-402550/37.	DR W71; 97-402550/37.
NPDB/V0198.			NPDB/V0198.	NPDB/V0198.	NPDB/V0198.
PT New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -			PT New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -	PT New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -	PT New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -
PR PR and related nucleic acid, useful for diagnosis and treatment of			PR PR and related nucleic acid, useful for diagnosis and treatment of	PR PR and related nucleic acid, useful for diagnosis and treatment of	PR PR and related nucleic acid, useful for diagnosis and treatment of
PS PS Claim 6; Fig 6; 136pp; French.			PS PS Claim 6; Fig 6; 136pp; French.	PS PS Claim 6; Fig 6; 136pp; French.	PS PS Claim 6; Fig 6; 136pp; French.
CC This is the amino acid sequence of the human protein SR-p70a. SR-p70a genes are transcription factors which may control the activity of p53-regulated genes, and are expressed by tumour suppressor genes related to the p53 gene family. The gene sequence was isolated from the human colon adenocarcinoma cell line HT-29, using primers V1506-7. The sequence can be used in the diagnosis and monitoring of cancer, especially CC neuroblastoma. The nucleic acid sequences and corresponding antisense CC sequences are also useful in gene therapy, e.g. to regulate apoptosis. Sequence 636 AA;			CC This is the amino acid sequence of the human protein SR-p70a. SR-p70a genes are transcription factors which may control the activity of p53-regulated genes, and are expressed by tumour suppressor genes related to the p53 gene family. The gene sequence was isolated from the human colon adenocarcinoma cell line HT-29, using primers V1506-7. The sequence can be used in the diagnosis and monitoring of cancer, especially CC neuroblastoma. The nucleic acid sequences and corresponding antisense CC sequences are also useful in gene therapy, e.g. to regulate apoptosis. Sequence 636 AA;	CC This is the amino acid sequence of the human protein SR-p70a. SR-p70a genes are transcription factors which may control the activity of p53-regulated genes, and are expressed by tumour suppressor genes related to the p53 gene family. The gene sequence was isolated from the human colon adenocarcinoma cell line HT-29, using primers V1506-7. The sequence can be used in the diagnosis and monitoring of cancer, especially CC neuroblastoma. The nucleic acid sequences and corresponding antisense CC sequences are also useful in gene therapy, e.g. to regulate apoptosis. Sequence 636 AA;	CC This is the amino acid sequence of the human protein SR-p70a. SR-p70a genes are transcription factors which may control the activity of p53-regulated genes, and are expressed by tumour suppressor genes related to the p53 gene family. The gene sequence was isolated from the human colon adenocarcinoma cell line HT-29, using primers V1506-7. The sequence can be used in the diagnosis and monitoring of cancer, especially CC neuroblastoma. The nucleic acid sequences and corresponding antisense CC sequences are also useful in gene therapy, e.g. to regulate apoptosis. Sequence 636 AA;
CC Best Local Similarity 100.0%; Pred. No. 1.1e-287; Mismatches 0; Indels 0; Gaps 0;			CC Best Local Similarity 100.0%; Pred. No. 1.1e-287; Mismatches 0; Indels 0; Gaps 0;	CC Best Local Similarity 100.0%; Pred. No. 1.1e-287; Mismatches 0; Indels 0; Gaps 0;	CC Best Local Similarity 100.0%; Pred. No. 1.1e-287; Mismatches 0; Indels 0; Gaps 0;
DB 1 MAQSTATSPDGTTFFHLWSSLEPDSTYFDLPLQSRRGNNNEVGGTDSMDVYLEGMTS 60			DB 1 MAQSTATSPDGTTFFHLWSSLEPDSTYFDLPLQSRRGNNNEVGGTDSMDVYLEGMTS 60	DB 1 MAQSTATSPDGTTFFHLWSSLEPDSTYFDLPLQSRRGNNNEVGGTDSMDVYLEGMTS 60	DB 1 MAQSTATSPDGTTFFHLWSSLEPDSTYFDLPLQSRRGNNNEVGGTDSMDVYLEGMTS 60
OY 1 YPGPHIFEVTFQQSSTAKSATWIVSPLLKKLYCQIAKICPIQKVSTTPPPGAIAMPV 180			OY 1 YPGPHIFEVTFQQSSTAKSATWIVSPLLKKLYCQIAKICPIQKVSTTPPPGAIAMPV 180	OY 1 YPGPHIFEVTFQQSSTAKSATWIVSPLLKKLYCQIAKICPIQKVSTTPPPGAIAMPV 180	OY 1 YPGPHIFEVTFQQSSTAKSATWIVSPLLKKLYCQIAKICPIQKVSTTPPPGAIAMPV 180
Db 1 YKAEAVTDVKRCNPFMNSCNCNSYRRPILITLLEMRDGGYLGRRSFEGRICAPGR 180			Db 1 YKAEAVTDVKRCNPFMNSCNCNSYRRPILITLLEMRDGGYLGRRSFEGRICAPGR 180	Db 1 YKAEAVTDVKRCNPFMNSCNCNSYRRPILITLLEMRDGGYLGRRSFEGRICAPGR 180	Db 1 YKAEAVTDVKRCNPFMNSCNCNSYRRPILITLLEMRDGGYLGRRSFEGRICAPGR 180
QY 61 VMAQENLSSMDQNSSSRAASASPTPEHAASVPTHSPYQAPSSTFDMSPAVIVPSNTD 120			QY 61 VMAQENLSSMDQNSSSRAASASPTPEHAASVPTHSPYQAPSSTFDMSPAVIVPSNTD 120	QY 61 VMAQENLSSMDQNSSSRAASASPTPEHAASVPTHSPYQAPSSTFDMSPAVIVPSNTD 120	QY 61 VMAQENLSSMDQNSSSRAASASPTPEHAASVPTHSPYQAPSSTFDMSPAVIVPSNTD 120
Db 61 VMAQENLSSMDQNSSSRAASASPTPEHAASVPTHSPYQAPSSTFDMSPAVIVPSNTD 120			Db 61 VMAQENLSSMDQNSSSRAASASPTPEHAASVPTHSPYQAPSSTFDMSPAVIVPSNTD 120	Db 61 VMAQENLSSMDQNSSSRAASASPTPEHAASVPTHSPYQAPSSTFDMSPAVIVPSNTD 120	Db 61 VMAQENLSSMDQNSSSRAASASPTPEHAASVPTHSPYQAPSSTFDMSPAVIVPSNTD 120
OY 121 YPGPHIFEVTFQQSSTAKSATWIVSPLLKKLYCQIAKICPIQKVSTTPPPGAIAMPV 180			OY 121 YPGPHIFEVTFQQSSTAKSATWIVSPLLKKLYCQIAKICPIQKVSTTPPPGAIAMPV 180	OY 121 YPGPHIFEVTFQQSSTAKSATWIVSPLLKKLYCQIAKICPIQKVSTTPPPGAIAMPV 180	OY 121 YPGPHIFEVTFQQSSTAKSATWIVSPLLKKLYCQIAKICPIQKVSTTPPPGAIAMPV 180
Db 121 YKAEAVTDVKRCNPFMNSCNCNSYRRPILITLLEMRDGGYLGRRSFEGRICAPGR 180			Db 121 YKAEAVTDVKRCNPFMNSCNCNSYRRPILITLLEMRDGGYLGRRSFEGRICAPGR 180	Db 121 YKAEAVTDVKRCNPFMNSCNCNSYRRPILITLLEMRDGGYLGRRSFEGRICAPGR 180	Db 121 YKAEAVTDVKRCNPFMNSCNCNSYRRPILITLLEMRDGGYLGRRSFEGRICAPGR 180
QY 181 EPQVGETFTILYNFMNCNSCNCNSYRRPILITLLEMRDGGYLGRRSFEGRICAPGR 300			QY 181 EPQVGETFTILYNFMNCNSCNCNSYRRPILITLLEMRDGGYLGRRSFEGRICAPGR 300	QY 181 EPQVGETFTILYNFMNCNSCNCNSYRRPILITLLEMRDGGYLGRRSFEGRICAPGR 300	QY 181 EPQVGETFTILYNFMNCNSCNCNSYRRPILITLLEMRDGGYLGRRSFEGRICAPGR 300
Db 241 EPQVGETFTILYNFMNCNSCNCNSYRRPILITLLEMRDGGYLGRRSFEGRICAPGR 300			Db 241 EPQVGETFTILYNFMNCNSCNCNSYRRPILITLLEMRDGGYLGRRSFEGRICAPGR 300	Db 241 EPQVGETFTILYNFMNCNSCNCNSYRRPILITLLEMRDGGYLGRRSFEGRICAPGR 300	Db 241 EPQVGETFTILYNFMNCNSCNCNSYRRPILITLLEMRDGGYLGRRSFEGRICAPGR 300

3.301	DRKADEHYREQQLQPLNNESSAKNGASKRAFKQSPAYPALGACYVKRRHDEDETYLQR	360
3.301	DRKADEHYREQQLQPLNNESSAKNGASKRAFKQSPAYPALGACYVKRRHDEDETYLQR	360
3.361	GRENFEILMKLKEELELMELYQPFLVDSYRQOQLLQRPISHQPPSTGPVLSPMNKYHGG	420
3.361	GRENFEILMKLKEELELMELYQPFLVDSYRQOQLLQRPISHQPPSTGPVLSPMNKYHGG	420
4.21	MNPKLPSVNLGVQPPHSSAAATPVLGPVPGMUNNHGAVPANGEMSSHSAAQMSVSSH	480
4.21	MNPKLPSVNLGVQPPHSSAAATPVLGPVPGMUNNHGAVPANGEMSSHSAAQMSVSSH	480
4.81	CTPPPPYHADPSLVSFLTGCGPCIEYFTSQGQSYTHQLNLTIEDLGALKIPEQTMT	540
4.81	CTPPPPYHADPSLVSFLTGCGPCIEYFTSQGQSYTHQLNLTIEDLGALKIPEQTMT	540
5.41	IWRQLDQKGHDYSTAQQLRSSNAATISIGGGELQRQVMEEAVHRVRTITIPNRG	600
5.41	IWRQLDQKGHDYSTAQQLRSSNAATISIGGGELQRQVMEEAVHRVRTITIPNRG	600
6.01	GPGGGPDEADFGDLPDKARKPIKEETEARIH	636
6.01	GPGGGPDEADFGDLPDKARKPIKEETEARIH	636

2

30651; standard; Protein; 635 AA.
30651; 7-APR-1993 (first entry)
human NBS-1 alpha Protein
NBS-1; p73; antibody; p53 responsive element; p53 promoter;
NBS-1; antibody; p53 dependent tumour; growth inhibition.
omo sapiens.
omo 350 kDa.
omo 100 kDa.

2-MAY-1997 / US 0464505.
2-MAY-1997 / US 0093516.
DAND) DANA FARBER CANCER INST INC.
POST C., Kaelin W.
CREATING SUBJECTS USING NBS-1 PROTEINS AND ANTIBODIES - used to
INTERACT WITH P53-RESPONSIVE GENES AND INHIBIT GROWTH OF
P53-DEPENDENT TUMOUR CELLS
EXAMPLE, Fig 1A; 65PP; English.

The method has been developed for treating a subject having a tumour comprising: (a) determining the protein level of NBS-1 Protein (also known as p53) expressed in the tumour cell and in a corresponding non-malignant cell; (b) selecting subjects having NBS-1 protein level comparable or below that in a corresponding normal cell; (c) elevating tumour cell NBS-1 level, where NBS-1 interacts with p53-responsive promoters. Also described in the invention is an antibody that specifically binds to NBS-1, and an antibody raised to the carboxy portion of NBS-1. NBS-1 can activate gene expression of p53-responsive genes and can inhibit cell growth in a p53-like manner. The present sequence represents the human NBS-1 protein from the present invention.

Sequence 635 AA; Identity Match 99.5%; Local Similarity 99.8%; Score 3367.5; DB 1; Pred. No. 3.1e-286; Length 635;

the conservative thesis 635; mismatches 0; indels 1;

1 MACSTATSPDGGSYFDLPOSSEHLSLEPDSNNEVGDSSMBVFLEGMT

THE JOURNAL OF CLIMATE

I. ANALYSIS OF THE INFLUENCE OF THE ENVIRONMENT ON THE DISEASE

61 YMAOFNLSSSTMDONSSRAASASPYTPEHAAASYPTHSPYAOPSSSTEDTMSPAVIPSNT

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61 VMAQFNLLSSTMQMSRAASASPYTPEHAASVPTHSPYAQPSSTSFTDTMSPAPIPNSNT

卷之三

121 YPGHHEEVIFQSSIAWIIISPLRNLLGIANICFIUNVSIRPICALKAM

Db	121	YPCGHFHEVTFQGSTAKSATWYSPPLKLYCOAKTCPIQIKYSTPPPGTATRAMPV	180
Qy	181	YKRAEHVTDVVKRCPNHELGDFNEGQSAPASHLIRVEGNNLSQYDDPTGROSYYVPP	240
Db	181	YKRAEHVTDVVKRCPNHELGDFNEGQSAPASHLIRVEGNNLSQYDDPTGROSYYVPP	240
Qy	241	EPPQGTETTILYNFMCNSSCVGGMNRPIIITLLEMRDGQVTRGRSRSGRICACPGR	300
Db	241	EPPQGTETTILYNFMCNSSCVGGMNRPIIITLLEMRDGQVTRGRSRSGRICACPGR	300
Qy	301	DRKADEDHYREQQALNESSAKNGAASKRAFKQSPPAVPAVLAGVYKRRHGEDDTYYLQVR	360
Db	301	DRKADEDHYREQQALNESSAKNGAASKRAFKQSPPAVPAVLAGVYKRRHGEDDTYYLQVR	360
Qy	361	GREENFEILMKLKESIEMELMVPQDPSYDSTRQQQQLQRPSHLOPSYGPVLSPMNVHGG	420
Db	361	GREENFEILMKLKESIEMELMVPQDPSYDSTRQQQQLQRPSHLOPSYGPVLSPMNVHGG	420
Qy	421	MNKPSVSYNOLVGQPPhSSAATPNPGVPGMNLNNHGHAYPANGEMSSHSQAQSMYSGSH	480
Db	421	MNKPSVSYNOLVGQPPhSSAATPNPGVPGMNLNNHGHAYPANGEMSSHSQAQSMYSGSH	480
Qy	481	CTPPPPHYADPSLVSLFTLGJCPNCIEYFTSQGLSIYHLQLNLTEIDLGALKIPEQYRT	540
Db	481	CTPPPPHYADPSLVSLFTLGJCPNCIEYFTSQGLSIYHLQLNLTEIDLGALKIPEQYRT	540
Qy	541	IWRGQDLQKGHDYSTAQQQLRSSNAATISIGGSEBLQRQRYMEAHTFRVHTTIPNRC	600
Db	541	IWRGQDLQKGHDYSTAQQQLRSSNAATISIGGSEBLQRQRYMEAHTFRVHTTIPNRC	600
Qy	601	GPGGGPDEWADEFGLPDCKARKQPIKEEETEEAIIH	636
Db	600	GPGGGPDEWADEFGLPDCKARKQPIKEEETEEAIIH	635
RESULT			
	3		
	W36182	W36182 standard; Protein: 637 AA.	
ID	W36182;		
AC	W36182;		
DT	27-APR-1998 (first entry)		
DE	Monkey P53 tumour suppressor-related protein SR-P70s.		
KW	SR-P70; monkey; transcription factor; p53; tumour suppressor gene;		
KW	homology; differential splicing; diagnosis; cancer; neuroblastoma;		
KW	gene therapy; apoptosis.		
OS	Cercopithecus aethiops.		
PN	WO97/28186A1.		
PD	07-AUG-1997.		
PF	03-FEB-1997; F00214.		
PR	02-FEB-1996; FR-0011309.		
PA	(SNFI) SANOFI SA.		
PI	Caput D, Ferrara P, Kaghad AM;		
DR	WPI: 97-422550/37.		
DR	N-PSDB; V01496.		
PT	New polypeptide(s) encoded by the SR-P70 tumour suppressor gene -		
PT	and related nucleic acid, useful for diagnosis and treatment of		
PT	tumours		
PS	Claim 1; Fig 4; 136pp. French.		
CC	This is the amino acid sequence of the protein SR-P70a from monkey ce		
CC	SR-P70 are transcription factors which may control the activity of		
CC	p53-regulated genes, and are expressed by tumour suppressor genes rel-		
CC	to the p53 gene family. The gene sequence was isolated from a cDNA		
CC	library by sequencing the inserts and comparing to sequence databases.		
CC	The protein sequence contains regions of homology to the p53 protein.		
CC	A second sequence (SR-P70b) was also isolated from the library and wa-		
CC	caused by differential splicing of the sequence (see V01497). The		
CC	sequences can be used in the diagnosis and monitoring of cancer,		
CC	especially neuroblastoma. The nucleic acid sequences and correspond-		
CC	ing antisense sequences, are also useful in gene therapy, e.g. to regulat-		
CC	apoptosis.		
SO	Sequence. 637 AA.		

RESULT 3 W36182

Query Match 97.7%; score 3304.5; DB 1; Length 637;
 Best Local Similarity 97.5%; Pred. No. 1e-280; Gaps 1;
 Matches 621; Conservative 4; Mismatches 11; Indels 1;

Qy 1 MAQSPATSPDGTTPEHLNLSSLEPPSTYFDLPOQSSRGNEVGGTDSMDVFLEGMTS 60
 Db 1 MAQSPATSPDGTTPEHLNLSSLEPPSTYFDLPOQSSRGNEVGGTDSMDVFLEGMTS 60

Qy 61 VMAQPNLSSMDQNSRAASAPSPVTPHEASVPHSPVQAPSSTEDMSPAVIPSTSND 120
 Db 61 VMAQPNLSSMDQNSRAASAPSPVTPHEASVPHSPVQAPSSTEDMSPAVIPSTSND 120

Qy 121 YPGPHFEVTFQQSSTAKSATWTSPLKLKYCQAKTCPIQIKVSTPPPGTAIRAMPV 180
 Db 121 YPGPHFEVTFQQSSTAKSATWTSPLKLKYCQAKTCPIQIKVSTPPPGTAIRAMPV 180

Qy 181 YKKAEHVTDTYKRCPNEHLGRDFNQGQASAPASHLJRVEGNLSQYDDPTGROSVVYP 240
 Db 181 YKKAEHVTDTYKRCPNEHLGRDFNQGQASAPASHLJRVEGNLSQYDDPTGROSVVYP 240

Qy 241 EPPQGTTEFTILYFMCSVCGMMNRPIIITLEFRDQYGLGRSFEGRICACGR 300
 Db 241 EPPQGTTEFTILYFMCSVCGMMNRPIIITLEFRDQYGLGRSFEGRICACGR 300

Qy 301 DRKADEDHYREQQALNESSAKNGAASKRAFKQSPPAVPAAGYKRRHGEDDETYLQVR 360
 Db 301 DRKADEDHYREQQALNESSAKNGAASKRAFKQSPPAVPAAGYKRRHGEDDETYLQVR 360

Qy 361 GRENEILMLKLESELMELYQPQYDTSRQQQOLQRPSPHSLQPSYGPVLSPMNKYHGG 420
 Db 361 GRENEILMLKLESELMELYQPQYDTSRQQQOLQRPSPHSLQPSYGPVLSPMNKYHGG 420

Qy 421 MNKLFPSYNOLYQOPPHSSATPNQGPYQPGMMLNNHGHAYPANGEMSSHSIQSMYSGSH 480
 Db 421 VNKLESVNQLYQOPPHSSATPNQGPYQPGMMLNNHGHAYPANSESSHTGTSQMSGSH 480

Qy 481 CTPPPPYADPSLVSFLTGIGCPNCIEYTSQGLQSIYLQNUTIEDGALKPEQYMT 540
 Db 481 CTPPPPYADPSLVSFLTGIGCPNCIEYTSQGLQSIYLQNUTIEDGALKPEQYMT 540

Qy 541 IWRGLQDLKGHDY STAQQLLRSSNAATISGSEGLORQTMEEAVFVRHTITIPNR 599
 Db 541 IWRGLQDLKGHDYAAQAQLLSSNAATISGSEGLORQTMEEAVFVRHTITIPNR 600

Qy 600 GGPGGGPDEADFDDLPDCDKARKOPIKEEFTEAIH 636
 Db 601 GGPAGGPDEADFDDLPDCDKARKOPIKEEFTEAIH 637

RESULT 4
 W36189 standard; Protein: 588 AA.
 ID W36189; PR-001309.
 AC W36189; (SNF1) SNOEI SA.
 DT 27-APR-1998 (first entry)
 DE Human P53 tumour suppressor-related protein SR-p70f.
 KW SR-p70; human; transcription factor; P53; tumour suppressor gene;
 KW homology; differential splicing; diagnosis; cancer; neuroblastoma;
 KW gene therapy; apoptosis.
 OS Homo sapiens.
 PN W0978186 AI.
 PD 07-AUG-1997.
 PP 03-FEB-1997; F00214.

PR 02-FBB-1996; FR-001309.
 PA Caput D, Ferrara P, Kaghad AM;
 DR WPI; 97-42550/37.
 DR N-PSDB; V01504.
 PT New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -
 PT and related nucleic acid, useful for diagnosis and treatment of
 PT tumours.
 Claim 1; Page 69-70; 136pp; French.
 PS This is the amino acid sequence of the human protein SR-p70f. SR-p70 are
 CC transcription factors which may control the activity of P53-regulated

genes, and are expressed by tumour suppressor genes related to the p53 gene family. The gene sequence was isolated from the human neuroblastoma cell line SK-N-SH, using primers V01515 and V01518. The SR-p70f gene sequence contains a 98 bp deletion between bases 24-25 as compared to the SR-p70a sequence (V01498). This deletion causes a loss of the downstream ATG (corresponding to an internal Met codon in SR-p70a). The protein is truncated by 48 amino acids at the N-terminus as compared to the SR-p70a protein (W36188). The sequence can be used in the diagnosis and monitoring of cancer, especially neuroblastoma. The nucleic acid sequences and corresponding antisense sequences, are also useful in gene therapy, e.g. to regulate apoptosis. Sequence 588 AA;

CC genes, and are expressed by tumour suppressor genes related to the p53 gene family. The gene sequence was isolated from the human neuroblastoma cell line SK-N-SH, using primers V01515 and V01518. The SR-p70f gene sequence contains a 98 bp deletion between bases 24-25 as compared to the SR-p70a sequence (V01498). This deletion causes a loss of the downstream ATG (corresponding to an internal Met codon in SR-p70a). The protein is truncated by 48 amino acids at the N-terminus as compared to the SR-p70a protein (W36188). The sequence can be used in the diagnosis and monitoring of cancer, especially neuroblastoma. The nucleic acid sequences and corresponding antisense sequences, are also useful in gene therapy, e.g. to regulate apoptosis. Sequence 588 AA;

Query Match 92.5%; Score 3130; DB 1; Length 588;
 Best Local Similarity 90.0%; Pred. No. 1.7e-265; Mismatches 0; Indels 0; Gaps 0;

Matches 588; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 49 MDYFHLEGMTISVMQAFNLSSTSMDQMSRAASASPYTEHAASVPTSPYAGPSSTDT 108
 Db 1 MDYFHLEGMTISVMQAFNLSSTSMDQMSRAASASPYTEHAASVPTSPYAGPSSTDT 60

Qy 109 MSAPAPVIPSNTDYPGPHFEVTFQOSSTAKSATTSYSPLLKLYCQIJKTCPIQKYSTP 168
 Db 61 MSAPAPVIPSNTDYPGPHFEVTFQOSSTAKSATTSYSPLLKLYCQIJKTCPIQKYSTP 120

Qy 169 PPGBTATRAMPVYKAERTYDVTYKRCPNEGOSAPASHLIRVEGNLSQLYVDD 228
 Db 121 PPGBTATRAMPVYKAERTYDVTYKRCPNEGOSAPASHLIRVEGNLSQLYVDD 180

Qy 229 PVTGROSVVYYEPPOVGTEFTILYNFMCNSCVGGMNRRPILLTLEMRGQVTLGRR 288
 Db 181 PVTGROSVVYYEPPOVGTEFTILYNFMCNSCVGGMNRRPILLTLEMRGQVTLGRR 240

Qy 289 SFEGRICACPGRDAKEDHYREQQALNESSAKNGAASKRAFKQSPPAVPAAGYKRRHGEDDETYLQVR 348
 Db 241 SFEGRICACPGRDAKEDHYREQQALNESSAKNGAASKRAFKQSPPAVPAAGYKRRHGEDDETYLQVR 300

Qy 349 HGDEDYYLQYGRENFELMLKLESLEMLYQPOLYDSYRQQLLQRPSHQLQPSY 408
 Db 301 HGDEDYYLQYGRENFELMLKLESLEMLYQPOLYDSYRQQLLQRPSHQLQPSY 360

Qy 409 PVLSPHANKYVGGMNRLPSYNQVQPPPHSSATAPNLGPVGPMLNNGHAYPANGEMSS 468
 Db 361 PVLSPHANKYVGGMNRLPSYNQVQPPPHSSATAPNLGPVGPMLNNGHAYPANGEMSS 420

Qy 469 SHSAQSMVYGSHTCPPPYHADPSLVSYFGLGCPCNIEYTSQGLQSIYLQNUTIEDGALKPEQYMT 528
 Db 421 SHSAQSMVYGSHTCPPPYHADPSLVSYFGLGCPCNIEYTSQGLQSIYLQNUTIEDGALKPEQYMT 480

Qy 529 GALKIPQEQYMTWGLQDLKGHDYSTAQQLLRSSNAATISGSEGLORQTMEEAVFVRHTITIPNR 588
 Db 481 GALKIPQEQYMTWGLQDLKGHDYSTAQQLLRSSNAATISGSEGLORQTMEEAVFVRHTITIPNR 540

Qy 589 RYRHITIPRGPGGGPDEADFGFDLPPDCDKARKQPKKEETTAETH 636
 Db 541 RYRHITIPRGPGGGPDEADFGFDLPPDCDKARKQPKKEETTAETH 588

RESULT 5
 W36187 standard; Protein: 587 AA.
 AC W36187; (first entry)
 DT 27-APR-1998
 DE Human P53 tumour suppressor-related protein SR-p70f.
 KW SR-p70; human; transcription factor; P53; tumour suppressor gene;
 KW homology; differential splicing; diagnosis; cancer; neuroblastoma;
 KW gene therapy; apoptosis.
 OS Homo sapiens.
 PN W0978186 AI.
 PD 07-AUG-1997; F00214.
 PP 03-FEB-1997; F00214.

'R 02-FEB-1996; FR-001309.
 'A (SFR) SANOFI SA;
 'I Caput D, Ferrara P, Raghad AM;
 'W WP1; 97-402550/37.
 'R N-PSDB; V01502.
 'T New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -
 'T and related nucleic acid, useful for diagnosis and treatment of
 'T tumours.
 'S Claim 1; Page 62-64; 136pp; French.
 'C This is the amino acid sequence of the human protein SR-p70d. R-p70 are
 'C transcription factors which may control the activity of p53-regulated
 'C genes, and are expressed by tumor suppressor genes related to the p53
 'C gene family. The gene sequence was isolated from the human colon
 'C neuroblastoma cell line IMR-32, using primers V015184-13. The SR-p70d
 'C protein sequence is 49 amino acids shorter with a divergence of the
 'C first 13 amino acids as compared to the SR-p70a protein (V015184).
 'C The sequence can be used in the diagnosis and monitoring of cancer,
 'C especially neuroblastoma. The nucleic acid sequences and corresponding
 'C antisense sequences, are also useful in gene therapy, e.g. to regulate
 'C apoptosis.
 'Q Sequence 587 AA;

Query Match 90.4%; Score 3058; DB 1; Length 587;

Best Local Similarity 100.0%; Pred. No. 3.3e-259; Mismatches 0;

Matches 574; Conservative 0; Indels 0; Gaps 0;

DE	Mouse p53 tumour suppressor-related protein SR-p70c.
KW	SR-p70; mouse; transcription factor; p53; tumour suppressor gene;
KW	homology; differential splicing; diagnosis; cancer; neuroblastoma;
KW	gene therapy; apoptosis.
OS	Mus musculus.
PN	W0728116-A1.
PD	07-AUG-1997.
PF	03-FEB-1997; F00214-309.
PR	02-FEB-1996; FR-001309.
PA	(SFR) SANOFI SA.
PI	Caput D, Ferrara P, Raghad AM;
DR	WP1; 97-402550/37.
DR	N-PSDB; V01502.
PT	New polypeptide(s) encoded by the SR-p70 tumour suppressor gene -
PT	and related nucleic acid, useful for diagnosis and treatment of
PT	tumours.
PS	Claim 1; Fig 7; 136pp; French.
CC	This is the amino acid sequence of the mouse protein SR-p70c. SR-p70
CC	are transcription factors which may control the activity of p53-regulated
CC	genes, and are expressed by tumour suppressor genes related to the p53
CC	gene family. The gene sequence was isolated from the mouse pituitary
CC	tumour cell line AtT 20, using primers V01508-9. The sequence can be
CC	used in the diagnosis and monitoring of cancer, especially neuroblastoma.
CC	The nucleic acid sequences and corresponding antisense sequences, are
CC	also useful in gene therapy, e.g. to regulate apoptosis.
SQ	Sequence 589 AA;

Query Match 83.6%; Score 2830.5; DB 1; Length 589;

Best Local Similarity 91.7%; Pred. No. 2.7e-239; Mismatches 19;

Matches 531; Conservative 19; Indels 7; Gaps 4;

Y	63 AOFNLISSTMDOMSSRAASASPYTPHEAASPYTHSPYAQSSTPDTMSAPVTSNTDYP 122
b	14 AOFNLISSTMDOMSSRAASASPYTPHEAASPYTHSPYAQSSTPDTMSAPVTSNTDYP 73
y	123 GPHFEFTVFOQSSTARATWVSPPLKKLYCQIAKTCPIQIKVSTPPGTATRAMPYK 182
b	74 GPHFEFTVFOQSSTARATWVSPPLKKLYCQIAKTCPIQIKVSTPPGTATRAMPYK 133
y	183 KAEHYTDVVKRCPNHELGRDFNEGQASAPASHLIRVEGNILSQTYDPTVGRQSYVVPYEP 242
b	134 KAEHYTDVVKRCPNHELGRDFNEGQASAPASHLIRVEGNILSQTYDPTVGRQSYVVPYEP 193
y	243 PQVGETFTILYNFMONNSCVGGMNRPLILITLEMRDQVLGRSFGRICACPGDR 302
b	194 PQVGETFTILYNFMONNSCVGGMNRPLILITLEMRDQVLGRSFGRICACPGDR 253
y	303 KADEDHYREQQALNESSAKNGASKRAFKOSPPAYPALGAGVKERRHGDEDTYLQVRQR 362
b	234 KADEDHYREQQALNESSAKNGASKRAFKOSPPAYPALGAGVKERRHGDEDTYLQVRQR 313
y	363 ENFEILMKLKESLELMELVQPLVDYSPQQQLLORPSHQOPSYGPVSPMKVHGMN 422
b	314 ENFEILMKLKESLELMELVQPLVDYSPQQQLLORPSHQOPSYGPVSPMKVHGMN 373
y	423 KLPVSNOVQGQPPhSSATPNLGPVPGMLNHRHAPANGENSSHQAQSAVSGSCT 482
b	374 KLPVSNCVLQGQPPhSSATPNLGPVPGMLNHRHAPANGENSSHQAQSAVSGSCT 433
y	483 PPPYHADPSLVSFLTGCGPCNCIEFTSQGLQSYTHLQNLTTEDLGALKIPDQYRTW 542
b	434 PPPYHADPSLVSFLTGCGPCNCIEFTSQGLQSYTHLQNLTTEDLGALKIPDQYRTW 493
y	543 RGLQDLKGHDYSTAQLRLSSNAATISIGGSGELQRORVMEAVHFRYRTHTIPNRGGP 602
b	494 RGLQDLKGHDYSTAQLRLSSNAATISIGGSGELQRORVMEAVHFRYRTHTIPNRGGP 553
y	603 GGGDEDAFDGFLPDKCKARQPIKEEFTAEIH 636
b	554 GGGDEDAFDGFLPDKCKARQPIKEEFTAEIH 587
ESULT	6
QY	541 IWRGLQDLKGHDYSTAQLRLSSNAATISIGGSGELQRORVMEAVHFRYRTHTIPNR 599
Db	36185 standard; Protein; 589 AA.
C	W3185;
C	27-APR-1998 (first entry)

Db	333	RGRERFEMRFELNEALELKD	352
	RESULT 14		
	R2238 ID R2238 standard; Protein: 393 AA.		
	AC R2238;		
	DT 23-JUL-1992 (first entry)		
	DE Sequence of 53 kd cellular protein.		
	KW Cancer therapy; cancer suppressor gene; oncogenesis.		
	OS Homo sapiens.		
	OS EP -7563-A.		
	PN 18-MAR-1992.		
	PP 23-AUG-1991; 307791.		
	PR 24-AUG-1990; US-513405.		
	PA (RECC) UNIV OF CALIFORNIA.		
	PI Lee WH, Chen PJ,		
	DR WPI; 92-09021/12.		
	N-PDB; Q22905.		
	PT Cloned P53 cDNA and protein prods. - for suppression of neoplastic phenotype e.g. in osteo-sarcoma(s), leukaemia(s), lymphoma(s), etc.		
	PT Claim 2; Page 14; 25pp; English.		
	PS P53 cDNA, or its gene prods., can be used to suppress and eradicate cancers caused by defective, mutant or absent cancer genes. Variant forms of p53 are found in human breast, lung or colon carcinoma, lymphoma, leukemia, etc., suggesting that mutation of the p53 genes is involved in oncogenesis. Specifically 273 Arg is replaced by 273 His, a mutation found exclusively in tumour cells.		
	CC Sequence 393 AA;		
	Query Match 21.4%; Score 724.5; DB 1; Length 393;		
	Best Local Similarity 41.0%; Prod. No. 2.8e-55; Mismatches 15%; Conservative 57; Indels 65; Gaps Matches 15;		
	QY 14 TFEHLWSSLEPDSTYFDLPOSSRGNNVEGGTDSNDVHLEMTTSYMAQFNLLSTMD 73		
	DB 18 TFSDLWKLPENNVLSPD-----SQAMDDMLMSPDIE 51		
	QY 74 QMSRAASASAPTYPEHAASVPTHSPYAQPSSTDTPAPAVP-----IPSNTDYGP 124		
	DB 52 QWFTEDPG ----- PDEARMPEAAPRVAAPAPAPTPA-----SQKTVQGS 106		
	QY 125 HHFETVFOSSSTAKSATWTPSPLKKLYCQIAKTCPOQIKVSTPPPGTCAIRAMPYKKKA 184		
	DB 107 YGFRLGEFLHSRCPHNRGLDFNEGOSAPASHLIRVEGNNSQYDVTGROSIVVYEPQQ 244		
	QY 185 EHVDYVKCPNPHLGRDFNEGOSAPASHLIRVEGNNSQYDVTGROSIVVYEPQQ 244		
	DB 167 QHMEEVRRCPHHERCSD SDG-LAPPOLIRVEGNLRLVEYLDRHTFRHSTVVPYEPPE 224		
	QY 245 VGETFTILYENMGNSSCVGGMNRPILLITLLEMRSQVLRGRSFEGRICACPGRDKA 304		
	DB 225 VGSQTTIHYNMNSSCVGGMNRPILTITLIDSSNLGNNSFEVRVACPGRDRT 284		
	QY 305 DEDHYREQALNESSAKNG-----AASKRKFQKSPPAVPLAGYKKKRHGDEDYY 356		
	DB 285 EEEUR-----KGKEPHHELPPSTKRALPNNTSSSP-----PKKPLDGEYFT 329		
	QY 357 LQVRGRNFETILMKLESLELME 379		
	DB 330 LQIRGRERFEMFRLNEALELK 352		
	RESULT 15		
	R26758 ID R26758 standard; peptide; 393 AA.		
	AC R26758;		
	DT 09-FEB-1993 (first entry)		
	P53 Point mutation; translocation; proto-oncogene; cancer;		

KW antigen-presenting cell; T-cell; HLA.
 OS Synthetic
 FH Key
 Location/Qualifiers
 FT misc_difference 273
 /label= "mutation"
 FT /note= "Arg -> any amino acid except Arg, see CC"
 FT GB2253211-A.
 PN PD SEP-1992.
 PD 26-FEB-1992; 004098.
 PR 26-FEB-1991; GB-003174.
 PA NHYD) NORSK HYDRO AS.
 PI Eriksen J, Gaardernck G, Geddedahl T;
 DR WPI: 92-284575/36.
 PT Synthetic oncogene protein peptide - used for treating and
 PT vaccinating against cancers
 PS Disclosure: Page 10; 78pp; English.
 CC New peptides, which have a point mutation or translocation compared to
 CC the correps. fragment of the proto-oncogene prod., correspond to,
 CC completely cover or are active fragments of a processed oncogene protein
 CC fragment as presented by a cancer cell or other antigen-presenting cell
 CC and are capable of inducing a specific T-cell response to the actual
 CC oncogene protein fragment as produced by the cell and processed and
 CC presented in the HLA mol.
 CC For example, a peptide fragment of p53 comprising at least
 CC mutations in position 2/3, in which position any amino acid except Arg
 CC may be located. The p53 sequence below was not disclosed in the
 CC specification, but retrieved by the indexer from Swiss-prot P04637.
 SQ

Query Match 21.4%; Score 724.5%; DB 1; Length 393;
 Best Local Similarity 42.1%; Freq. No. 2.8e-55;
 Matches 160; Conservative 58; Mismatches 103; Indels 59; Gaps 10;

Qy 14 TFEHLWSSLRFDSTYFDLPOSSRGNEVYGGTDDSSMDVFHLEGMTTSYMAOFNLSSSTMID 73
 Db 18 TFSDIWKLPLPENVYLSPLP-SQAMDDLMUSPD-----TEQWTEDEDPPD 61

Qy 74 QMSSRAASASPYTPHEAASYPHTSFYQAQSSTFTMSPLP-----VAPSNTDYPGPPHF 127
 Db 62 EAIPRPEAAFPVAPAAFP-----AAP-----APAPSWPLSSPSQKTYQGSGF 109

Qy 128 EVTFQOSSTAKSATVYSPILKLYCQIAKTCPIQIKYSTPPPGTAIRAMPYKAHV 187
 Db 110 RLGFHLHSCTAKSVCITYSPALNKMFQCOLAKTCPIQYDPTVTRAMAYKQSQHM 169

Qy 188 TDVYRCPNIELGRDNFEGGASAPASHLIREVEGNNSQYVYDDPVYGRQSYVVPPQVGT 247
 Db 170 TEVYRCRPHERCSD-SDG-LAPPOLIRVEGLNRYEILDRTNRHSVVPYEPPEVGS 227

Qy 248 EFTTLIYNFWCNSSCGGAGNRRPTLTLEMRDQVGRSRSGRIGACPGIDRRADED 307
 Db 228 DCTTHYNYCNSSCGGAGNRRPTLTLEDSSNLGRNSFVRVACPGDRRTBEE 287

Qy 308 HYREQOALNESSSAKING-----ASKRAFKQSPSPAYPALGAGVKRRHGDDTYYLQV 359
 Db 288 NLR-----KGEPFHIELPPOSTKRALPNNTSSSPQ-----PKKKPLDGEYFTIQI 332

Qy 360 RGRENFEILMKLKESELME 379
 Db 333 RGREFEMFRELNEALELK 352